OBJECTIVES: The Senior Medical Director leads and drives strategy for the overall clinical development (taking into consideration the medical, scientific, regulatory and commercial issues) of assigned Bolt Biotherapeutics pipeline compounds. Leads a multi-disciplinary, matrix team through complex decisions. This individual has the ultimate responsibility to inform development decisions assessing and integrating input from various disciplines to create, maintain, and execute a U.S. and global clinical development plan that will result in the regulatory approval of the assigned compound in multiple regions as well as inform the value dossiers required for reimbursement. Applies clinical/medical decision-making to clinical development issues.

ACCOUNTABILITIES:
- Clinical Development team participation and leadership
- Synopsis / Protocol Development, Study Execution, & Study Interpretation
- Trial Medical Monitoring
- Key Stakeholder development
- Due Diligence, Business Development and Alliance Projects
- Hires, manages, mentors, motivates, empowers, develops and retains staff to support assigned activities.

EDUCATION, EXPERIENCE, KNOWLEDGE AND SKILLS:
- MD required with 5 years of clinical research experience within the academic and/or pharmaceutical industry (e.g., CRO health-related consulting company, or biomedical/clinical experience.)
- Previous experience successfully leading a clinical development team/matrix team with responsibility for studies in multiple regions preferred.
- IND submission experience preferred.
- Ability to proactively predict issues and solve problems
- Ability to inform and drive decision-making within a multi-disciplinary team
- Diplomacy and positive influencing abilities
- Oncology therapeutic area knowledge relevant to mechanism of action preferred

TRAVEL REQUIREMENTS:
- Ability to drive to or fly to various meetings or client sites, including overnight trips. Some international travel may be required.
- Requires approximately 15 - 25% travel.